 INFORMATION DISCLOSURE CITATION Form PTO-1449 (Modified) <i>(Use several sheets if necessary)</i>	ATTY. DOCKET NO. UCAL161DIV	SERIAL NO. 09/922,483
	APPLICANT Finkbeiner, et al.	
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U.S. PATENT DOCUMENTS

Examiner Initial	Document Number	Date	Name	Class	Subclass	Filing Date If Appropriate

FOREIGN PATENT DOCUMENTS

Document Number	Date	Country	Class	Subclass	Translation
AA WO 97/17445	15-May-97	PCT	—	—	Yes No

OTHER ART (Including Author, Title, Date, Pertinent Pages, Etc.)

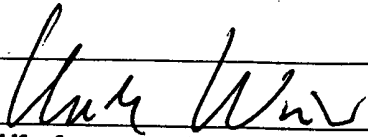
AB	Heiser et al. (2000) "Inhibition of huntingtin fibrillogenesis by specific antibodies and small molecules: Implications for Huntington's disease therapy." <i>PNAS</i> , Vol. 97(12):6739-6744.
AC	Imbert, et al., "Cloning of the Gene for Spinocerebellar Ataxia 2 Reveals a Locus With High Sensitivity to Expanded CAG/glutamine Repeats", <i>Nat. Genet</i> (Nov. 1996) 14:285-291
AD	Johansson et al. "Liver cell uptake and degradation of soluble immunoglobulin G immune complexes in vivo and in vitro in rats", <i>Hepatology</i> , Vol. 24 (1996) pp.169-175.
AE	Lunkes, et al., "Properties of Polyglutamine Expansion in Vitro and in Cellular Model for Huntington's Disease", <i>Phil. Trans. R. Soc. Lond. B</i> (1999) 354:1013-1019
AF	Persichetti et al. (1996) "Differential Expression of Normal and Mutant Huntington's Disease Gene Alleles." <i>Neurobiology of Disease</i> , Vol. 3:183-190.
AG	Persichetti, et al., "Mutant Huntingtin Forms in Vivo Complexes With Distinct Context-Dependent Conformations of the Polyglutamine Segment", <i>Neurobiology of Disease</i> (1999) 6:364-375
AH	Stevanin, et al., "Screening for Proteins With Polyglutamine Expansions in Autosomal Dominant Cerebellar Ataxias", <i>Hum. Mol. Genet.</i> (Dec. 1996) 5:887-1892
AI	Takeuchi, et al., "Molecular Cloning and Expression of a Novel Human cDNA Containing CAG Repeats", <i>Gene</i> (1997) 204:71-77
AJ	Tromier, et al., "Polyglutamine Expansion as a Pathological Epitope in Huntington's Disease and Four Dominant Cerebellar Ataxias", <i>Nature Genet.</i> (1995) Pages 403-406
AK	Trottier, et al., "Cellular Localization of the Huntington's Disease Protein and Discrimination of the Normal and Mutated Form", <i>Nature Genetics</i> (May 1995) 10:104-110

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